

What is claimed is:

- 1. Synthetic analogues of PtdIns(4,5)P₂ incomporating one or more of the following modifying structural features, (i) the 2-OH is rendered non-nucleophilic by: (a) derivatization, exemplified by 2-OCOR of 2-OR, R = alkyl, substituted alkyl or alkenyl, particularly 2-OAC, or by replacement in 2-deoxyhalo, and 2-dideoxyhalo, particularly 2-O-deoxyfluoro, (ii) photoaffinity, fluorescent, spin, other reporter groups, and conjugands for linking to polymer, chromatographic matrix, or gold surfaces are incorporated in the fatty acyl or inositol residues as exemplified in structure 2 wherein the absolute stereochemistry of the natural PtdIns(4,5)P₂ is maintained.
- 2. Analogues of Claim 1 wherein the glycerol esters are replaced by ether bonds.
- 3. Analogues of Claims 1 and 2 applied as analytical reference and research reagents.
- 4. A multi-step method for the synthesis of 2-modified PtdIns(4,5)P₂ analogues as described in FIG. 1 and 2.
- 5. 1D-3,6-Di-O-benzyl-4,5-O-cycloHexylidene-myo-inositol as a starting material for the synthesis of Claim 3.
- 6. Analogues of 1D-3,6-di-O-benzyl-445-O-cyclohexylidene-myo-inositol wherein the 4,5-O-protection is derived from acetonel benzaldehyde, camphor or equivalent in place of cyclohexanone.
- 7. Analogues of 1D-3,6-di-O-benzyl-4,5-O-cyclohexylidene-myo-inositol wherein the 3,6-O-protection is provided by allyl, 1-propenyl, p-methoxybenzyl, and equivalent groups removable by reagents other than catalytic hydrogenolysis.
- 8. Key phosphodiester intermediates in synthesis according to Claims 3 and 4 formed by the reaction of a selectively protected myo-inositol and an sn-3-phosphatidic acid or glycero-ether analogues.
- 9. A method for synthesis of $PtdIns(4,5)P_2$ analogues based on derivatization by acylation, phosphorylation and equivalent reactions with a selectively protected $PtdIns(4,5)P_2$ carrying a free hydroxyl in the inositol.
- 10. Phosphoinositide analogues modified as in Claims 1 and 2 but based on a PtdIns-mono or poly-phosphate other than PtdIns(4,5)P₂.

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